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				ART UNIT	PAPER NUMBER
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			DATE MA	ILED:	
		examiner interview su	mmary record		
All participants (applican	t, applicant's representa	tive, PTO personnel):			
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(1) John Store	(Appi Rep)	(3)	refer senbert	(Inventur	3
(2) Jean Duva	all (Appi-Rep)	(3)	PATERIA DURY	CELAMINE	R)
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			renrecentative)		
•		en to ロ applicant ロ applicant's Yes 없 No. If yes, brief description			
Exhibit shown or demon	Stration Conducted.	165 Q 140. II yes, biloi description	10 4		
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Agreem of Dwas rea	ched with respect to son	ne or all of the claims in question.	স was not reached.		
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Claims discussed:	All pending			·	
ld ntification of prior art	discussed:	NE			
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D scription of the gener	al nature of what was ag	preed to if an agreement was reache	d, or any other comments: _		
-	500	attached for spec	fic.		
		sed enablements			•
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<u>35USC</u>	. 112 1 st par	agraph.			
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(A fuller description if pe	acceptant and a copy of	the amendments, if available, which	the examiner agreed would	render the claims	allowable must b
attached. Also, where n	o copy of the amendme	nts which would render the claims a	llowable is available, a sumn	nary thereof must	be attached.)
₩1. It is not necessa	ary for applicant to provid	de a separate record of the substanc	e of the interview.		
WAIVED AND MUST IN	ICLUDĖ THE SUBSTAN	to indicate to the contrary, A FORMA ICE OF THE INTERVIEW (e.g., item given one month from this interview of	ns 1-7 on the reverse side of	this form). If a re-	sponse to the last Office
 2. Since the example requirements the response requirements. 	nin r's int rview summar nat may be pres nt in the rements of the last Office	y ab v (including any attachments e last Office action, and since the cla e action. Applicant is not r lieved for) reflects a complete respons tims are now allowable, this	se to each of the c	objections, rejections and consider d to fulfill th
box 1 above is	also checked.		. 0	_	

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Interview Summary 12 December 1996

Discussion centered on five basic points.

- 1. The differences between Alzheimer's Disease and the transgeneic animal models of beta-amyloidosis. Discussed the fine point in regard to the lack of neurofibrillary tangles with respect to any existing animal or animal model. Discussed restriction of claims to potentially beta-amyloid deposition or secreting models. Support for this limitation in specification was queried by the examiner. Concurred that the transgeneic animal models were a model for beta-amyloidosis.
- 2. The ability of the assay to detect smaller samples which would be required in small rodents. Discussed that the claim language requires preadministration sampling, administration of the substance and postadministration sampling in the sample. A temporal sequence is implied by the antecedent basis of the claim with regard to "the non-human animal". Applicants were going to consider both points and potentially supply data showing that the assay is sensitive enough to assay smaller samples than the 100 ul used for human CSF in the specification. Applicants were also providing evidence that the assay can distinguish the transgene from the endogenous peptide in the transgeneic models of beta-amyloidosis.
- 3. Enablement regarding transgeneic animals will be supported by evidence (i.e. scientific papers to be provided).
- 4. Evidence regarding the establishment of more than one assay to determine the presence of the AB(x-≥41) commensurate in scope with the assay.
- 5. Arguments will be presented regarding screening nature of the assay rather than therapeutic with regards to Alzheimer's disease.

Exr. Patricia A. Duffy, Ph.D.

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